Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application.

Listing of Claims:

Claims 1-155 (cancelled)

Claim 156 (currently amended):

A compound of the formula

or <u>a</u> its pharmaceutically acceptable salt thereof.

Claims 157-162 (cancelled)

Claim 163 (currently amended):

A pharmaceutical composition comprising a compound of

the formula

or <u>a</u> its pharmaceutically acceptable salt thereof, in combination with β -L-deoxyribothymidine.

Claims 164-174 (cancelled)

Claim 175 (currently amended): A compound of the formula:

$$\begin{array}{c} NR^3R^4 \\ N \\ N \\ O \end{array}$$

or a its pharmaceutically acceptable salt thereof, wherein

R² is selected from the group consisting of straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, CO-aryloxyalkyl, CO-substituted aryl, alkylsulfonyl, arylsulfonyl, aralkylsulfonyl, an amino acid residue, mono, di, or triphosphate, or a phosphate prodrug; and

R³ and R⁴ are independently H, straight chained, branched or cyclic alkyl, dialkylaminoalkylene, CO-alkyl, CO-aryl, CO-alkoxyalkyl, CO-aryloxyalkyl, CO-substituted aryl, alkylsulfonyl, arylsulfonyl, aralkylsulfonyl, amino acid residue, mono, di, or triphosphate, or a phosphate prodrug.

Claim 176 (cancelled)

Claim 177 (cancelled)

Claim 178 (cancelled)

Claim 179 (currently amended): The compound of claim 175, wherein R^2 is an amino acid residue of the formula $C(O)C(R^8)(R^9)(NR^{10}R^{11})$, wherein

R⁸ is the side chain of an amino acid and wherein R⁸ can optionally be attached to R¹⁰ to form a ring structure; or alternatively, R⁸ is an alkyl, aryl, heteroaryl or <u>heterocycle</u> heterocyclic moiety;

R⁹ is hydrogen, alkyl or aryl; and

R¹⁰ and R¹¹ are independently hydrogen, acyl or alkyl.

Claim 180 (previously presented): The compound of claim 179, wherein R² is L-valinyl.

Claim 181 (previously presented): The compound of claim 175, wherein R³ and R⁴ are hydrogen.

Claim 182 (previously presented): The compound of claim 175, wherein R^3 is hydrogen and R^4 is dimethylaminomethylene.

Claim 183 (previously presented): The compound of claim 175, wherein R^3 is hydrogen and R^4 is CO-alkyl.

Claim 184 (previously presented): The compound of claim 175, wherein R^3 is hydrogen and R^4 is CO-methyl.

Claim 185 (previously presented): The compound of claim 175, wherein R^3 is hydrogen and R^4 is L-valinyl.

Claim 186 (currently amended): A compound of the formula:

or a its pharmaceutically acceptable salt thereof, wherein

R² is selected from the group consisting of straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, CO-aryloxyalkyl, CO substituted aryl, alkylsulfonyl, arylsulfonyl, aralkylsulfonyl, an amino acid residue, mono, di, or triphosphate, or a phosphate prodrug.

Claim 187 (cancelled)

Claim 188 (cancelled)

Claim 189 (cancelled)

Claim 190 (currently amended): The compound of claim 186, wherein \mathbb{R}^2 is an amino acid residue of the formula $C(O)C(\mathbb{R}^8)(\mathbb{R}^9)(\mathbb{NR}^{10}\mathbb{R}^{11})$, wherein

R⁸ is the side chain of an amino acid and wherein R⁸ can optionally be attached to R¹⁰ to form a ring structure; or alternatively, R⁸ is an alkyl, aryl, heteroaryl or <u>heterocycle</u> heterocyclic moiety;

R⁹ is hydrogen, alkyl or aryl; and

R¹⁰ and R¹¹ are independently hydrogen, acyl or alkyl.

Claim 191 (previously presented): The compound of claim 190, wherein R² is L-valinyl.

Claim 192 (currently amended): A pharmaceutical composition comprising an effective anti-HBV amount of a compound of the formula:

or a its pharmaceutically acceptable salt thereof, wherein

R² is selected from the group consisting of straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, CO-aryloxyalkyl, CO-substituted aryl, alkylsulfonyl, arylsulfonyl, aralkylsulfonyl, an amino acid residue, mono, di, or triphosphate, or a phosphate prodrug; and

R³ and R⁴ are independently H, straight chained, branched or cyclic alkyl, dialkylaminoalkylene (in particular, dimethylaminomethylene), CO-alkyl, CO-aryl, CO-alkoxyalkyl, CO-aryloxyalkyl, CO-substituted aryl, alkylsulfonyl, arylsulfonyl, aralkylsulfonyl, amino acid residue, mono, di, or triphosphate, or a phosphate prodrug; with a pharmaceutically acceptable carrier or diluent.

Claim 193 (cancelled)

Claim 194 (cancelled)

Claim 195 (cancelled)

Claim 196 (currently amended): The pharmaceutical composition of claim 192, wherein R² is an amino acid residue of the formula C(O)C(R⁸)(R⁹)(NR¹⁰R¹¹), wherein R⁸ is the side chain of an amino acid and wherein R⁸ can optionally be attached to R¹⁰ to form a ring structure; or alternatively, R⁸ is an alkyl, aryl, heteroaryl or heterocycle heterocyclic moiety;

R⁹ is hydrogen, alkyl or aryl; and

 R^{10} and R^{11} are independently hydrogen, acyl or alkyl.

Claim 197 (previously presented): The pharmaceutical composition of claim 196, wherein R² is L-valinyl.

Claim 198 (previously presented): The pharmaceutical composition of claim 192, wherein R³ and R⁴ are hydrogen.

Claim 199 (previously presented): The pharmaceutical composition of claim 192, wherein R³ is hydrogen and R⁴ is dimethylaminomethylene.

Claim 200 (previously presented): The pharmaceutical composition of claim 192, wherein R³ is hydrogen and R⁴ is CO-alkyl.

Claim 201 (previously presented): The pharmaceutical composition of claim 192, wherein R³ is hydrogen and R⁴ is CO-methyl.

Claim 202 (previously presented): The pharmaceutical composition of claim 192, wherein R³ is hydrogen and R⁴ is L-valinyl.

Claim 203 (currently amended): A pharmaceutical composition comprising an effective anti-HBV amount of a compound of the formula:

or a its pharmaceutically acceptable salt thereof, wherein

R² is selected from the group consisting of straight chained, branched or cyclic alkyl, CO-alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, CO-aryloxyalkyl, CO substituted aryl, alkylsulfonyl, arylsulfonyl, aralkylsulfonyl, an amino acid residue, mono, di, or triphosphate, or a phosphate prodrug;

with a pharmaceutically acceptable carrier or diluent.

Claim 204 (cancelled)

Claim 205 (cancelled)

Claim 206 (cancelled)

Claim 207 (currently amended): The pharmaceutical composition of claim 203, wherein R^2 is an amino acid residue of the formula $C(O)C(R^8)(R^9)(NR^{10}R^{11})$, wherein R^8 is the side chain of an amino acid and wherein, as in proline, R^8 can optionally be attached to R^{10} to form a ring structure; or alternatively, R^8 is an alkyl, aryl, heteroaryl or

heterocycle heterocyclic moiety; R⁹ is hydrogen, alkyl or aryl; and

 R^{10} and R^{11} are independently hydrogen, acyl or alkyl.

Claim 208 (previously presented): The pharmaceutical composition of claim 207, wherein R² is L-valinyl.

Claim 209 (currently amended): A pharmaceutical composition comprising a compound of the formula

or <u>a</u> its pharmaceutically acceptable salt thereof, with a pharmaceutically acceptable carrier or diluent.

Claims 210-241 (cancelled)